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APPLICATION NO. FILING DATE		FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
09:632,812	C	08:04/2000	Steven H. Coberly	9323.00001 2522	
22907	7590	03.20.2003			
BANNER &	witco)FF	EXAMINER		
1001 G STREET N W SUITE 1100				BARRY, CHESTER T	
WASHINGTON, DC 20001			ART UNIT	PAPER NUMBER	
				1724	11
				DATE MAILED: 03/20/2003	111

Please find below and/or attached an Office communication concerning this application or proceeding.

^ ′	Application No.	Applicant(s)					
	09/632,812	COBERLY ET AL.					
Office Action Summary	Examiner	Art Unit					
•	Chester T. Barry	1724					
The MAILING DATE of this communication ap	· •						
Period for Reply							
A SHORTENED STATUTORY PERIOD FOR REPL THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1 after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a replant of the provision of the provi	136(a). In no event, however, may a reply be to only within the statutory minimum of thirty (30) da will apply and will expire SIX (6) MONTHS from e. cause the application to become ABANDON	imely filed ays will be considered timely in the mailing date of this communication ED (35 U S C § 133)					
Status	Documbar 2002						
1) Responsive to communication(s) filed on <u>18</u> 2a) This action is FINAL . 2b) This action is FINAL .	his action is non-final.						
, <u> </u>		prosecution as to the merits is					
closed in accordance with the practice under	r Ex parte Quayle, 1935 C.D. 11,	453 O.G. 213.					
Disposition of Claims							
4) Claim(s) 1-12 is/are pending in the applicatio							
	4a) Of the above claim(s) is/are withdrawn from consideration.						
	Claim(s) is/are allowed.						
	6) Claim(s) 1-12 is/are rejected.						
7) Claim(s) is/are objected to.							
8) Claim(s) are subject to restriction and/o	or election requirement.						
9) The specification is objected to by the Examine	er						
10) The drawing(s) filed on is/are: a) acce		aminer.					
Applicant may not request that any objection to the							
11) The proposed drawing correction filed on is: a) approved b) disapproved by the Examiner.							
If approved, corrected drawings are required in re	eply to this Office action.						
12) ☐ The oath or declaration is objected to by the E	xaminer.						
Priority under 35 U.S.C. §§ 119 and 120							
13) Acknowledgment is made of a claim for foreig	gn priority under 35 U.S.C. § 119	(a)-(d) or (f).					
a) ☐ All b) ☐ Some * c) ☐ None of:							
1. Certified copies of the priority documen	its have been received.						
2. Certified copies of the priority documen	2. Certified copies of the priority documents have been received in Application No						
3. Copies of the certified copies of the pricapplication from the International B* See the attached detailed Office action for a lis	ureau (PCT Rule 17.2(a)).						
14) Acknowledgment is made of a claim for domes	tic priority under 35 U.S.C. § 119	e(e) (to a provisional application).					
 a) ☐ The translation of the foreign language pr 15) ☐ Acknowledgment is made of a claim for domes 							
Attachment(s)							
 Notice of References Cited (PTO-892) Notice of Draftsperson's Patent Drawing Review (PTO-948) Information Disclosure Statement(s) (PTO-1449) Paper No(s) 	5) Notice of Informa	ary (PTO-413) Paper No(s) al Patent Application (PTO-152)					

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Claims 9 - 12 are rejected under §112, first paragraph, for failure of the application as originally filed to describe the claims invention, and under §251 for addition of new matter to the application.

Each of claims 9 – 12 requires that "substantially all heat stable anions transferred to the resin in step (a)" are removed by the alkali metal hydroxide added in step (b) to regenerate the resin. There is no evidence that applicants were in possession of exchanging hydroxide anions for only those anions that were transferred to the resin in the preceding step. While there is support for the concept of regenerating the capacity of the resin to substantially the same capacity as that of the previous cycle, there is no showing that applicant ever identified that the sites regenerated by hydroxide were the same sites to which anions were transferred during the previous exhaustion cycle.

USP 6245128 to George (ostensibly assigned to Mobil Oil Corporation) is cited of interest for what it claims rather than for what it teaches. Note in particular it recital in the claims of sodium chloride solution as a regenerant. Note also USP 4267159 to Crits for the teaching that a strong alkali regenerant for a Type II resin can be the salt of a strong base, such as sodium chloride, or the strong base itself, e.g., sodium hydroxide. Crits col 3 lines 30-38.

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Crits is also cited of interest for the distinction it makes between Type I and Type II resins. According to Crits, a Type I resin has trimethylamine functional groups whereas a Type II resin relies on dimethyl ethanolamine for functionality.

Crits (example 2) is also cited of interest for a cyclic process in which in the ninth run, a Type II resin is exhausted and then partially regenerated to a capacity of 0.7 kilograin per cubic foot using *calcium* hydroxide solution (not an alkali metal hydroxide). Thereafter, the salt of a strong base, i.e., an 15% sodium chloride solution, is used to regenerate the capacity of the Type II resin to within about 15% of its original virgin capacity ((2.1-1.8)/1.8x100%). Crits' does not describe regenerating in a subsequent step the very same sites to which anion were transferred in the next preceding step. Critis' regeneration to within 15% of the original capacity using the salt of a strong base did not restore the capacity of the resin to "substantially [the same]" capacity as that resulting from the previous regeneration cycle. It is also noted that in the 9 runs of exhaustion and regeneration using weak base, ie., calcium hydroxide, the capacity decreased from 2.1 to 0.7 kilograins per cu. ft. If the extent of each regeneration were consistent one run to the next, the extent of regeneration per pass or cycle would have been about 88.5%, or approximately to within only about 11.5% of the capacity of the preceding cycle. If, on the other hand, strong base were used, e.g., 0.1 – 0.3 M NaOH solution, one of skill would have expected each pass to exceed about 98.3% of the previous capacity, or to within about 1.7% of the previous cycle's capacity, or to

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substantially the same capacity as in the next previous cycle $(2.1*(1-0.017)^9 = 1.8)$. Accordingly, one of skill would have expected a single regeneration cycle using strong base, such as 15% sodium hydroxide, to regenerate a Type II resin exhausted with heat stable salts to substantially the same capacity as that which resulted from the previous regeneration cycle.

Claims 1 - 12 are rejected under §112, 2nd paragraph, for failing to particularly point out and distinctly claim the subject matter for which patent protection is sought. Each of claims 1 – 12 recites a "Type II strong base anion exchange resin," but the specification does not describe what a Type II resin is. Claims are interpreted as the person of ordinary skill would have understood than at the time the invention was made. Without an adequate description of what applicants regard as a Type II resin, the skilled art artisan must rely on the teachings of the prior art to come to grips with what patentees must have had in mind when they adopted the "Type II" terminology to define their invention. Importantly, applicants on several occasions have distinguished their inventions from similar processes using Type I resins.

In light of the prior art, the skilled artisan is not put on fair notice of reasonably precise limits of the claim invention: Crits suggests that the feature distinguishing Type II from Type II resins is reliance on dimethyl ethanolamine and trimethyamine groups for anion exchange functionality, respectively. See also USP 5846883 Boone in this regard, too. USP 5692461 to Crovato, on the other hand, suggests that the difference

¹ In applicant's specification, the largest disparity between the capacity of one run and the subsequent run was 51%

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between a "Type 1" and a "Type 2" strong base anion exchange resin is the identity of the anion attached to the regenerated resin: Type 1 resins include chloride anions on trimethylamine functional groups whereas Type 2 resins include hydroxide anions on trimethylamine functional groups. See Crovato at col 25 lines 38-61. Perry's Chemical Engineering Handbook defines it not in terms of chemical constituents or in terms of the identity of the form of the resin (chloride form or hydroxide form), but rather in terms of the shape of the two component exchange isotherms. See Perry Fig. 16 - 8. In light of these various very different notions of what a "Type II" anion exchange resin is, the skilled artisan is hard-pressed to discern what applicants' invention is given his failure to define what he meant by it as his own lexicographer.

Perry is also cited for the recognition in this art of the concept of cyclic steady state. See page 16-38. USP 5254368 to Kadlec is cited for it discussion of reaching a cyclic steady state (col 6 line 45-52). USP 4344851 to Sherman is cited for the discussion of cyclic steady state in the context of an ion exchange process. USP 4768345 to Kardas and USP 4363327 to Clark are also cited of interest with regard to the understandings in various allied fields of the concept of cyclic steady state.

 $\mathsf{Respectfully},$

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